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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 2

Complete if Known	
Application Number	09/863,049
Filing Date	May 22, 2001
First Named Inventor	Sue J. Kenwrick
Art Unit	N/A
Examiner Name	Not Yet Assigned
Attorney Docket Number	HO-P01961US1

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number: Number-Kind Code[2] *(if known)*	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document: Country Code[3]-Number[4]-Kind Code[5] *(if known)*	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]

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NON PATENT LITERATURE DOCUMENTS

Examiner Initials	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
	CA	Mutations in Brief, Human Mutation Vol. 12, No. 5, (October 7, 1998), pp. 361-363.	
	CB	Rothnie, Helen, M. et al.; Polyadenylation in Rice Tungro Bacilliform Virus: cis-Acting Signals and Regulation, Journal of Virology (May 2001), Vol. 75, No. 9, pp. 4184-4194.	
	CC	McQueen, Karina L., et al.; Functional analysis of 5' and 3' regions of the closely related Ly49c and j genes, Immunogenetics (2001) Vol. 52, pp. 212-223.	
	CD	Graber, Joel H., et al.; In silico detection of control signals: mRNA 3' -end-processing sequences in diverse species, PNAS (November 23, 1999), Vol. 96, No. 24, pp. 14055-14060.	
	CE	Xu, Zhi-Li, et al.; Optimization of transcriptional regulatory elements for constructing plasmid vectors, Int'l Journal of Genes and Genomes, (2001), Vol. 272, pp. 149-156.	
	CF	Shamsher, Monee K., et al.; Identification of an intronic regulatory element in the human protein C (PROC) gene, Human Genetics (2000), Vol. 107, pp. 458-465.	
	CG	Ionasescu, V.V., et al.; Mutations of the noncoding region of the connexin32 gene in X-linked dominant Charcot-Marie-Tooth neuropathy, American Academy of Neurology (1996), Vol. 47, pp. 541-544.	
	CH	Harland, Mark, et al.; Mutation Screening of the CDKN2A Promoter in Melanoma Families, Genes, Chromosomes & Cancer (2000) Vol. 28, pp. 45-57.	
	CI	Pyne, Michael T., et al.; The BRCA2 genetic variant IVS7+ 2T - G is a mutation, J. Hum. Genetics (2000), Vol. 45, pp. 351-357.	
	CJ	Jannssen, R.J., et al.; A branch site mutation leading to aberrant splicing of the human tyrosine hydroxylase gene in a child with a severe extrapyramidal movement disorder, Am. Hum. Genet. (2000), Vol. 64, pp. 375-382.	
	CK	Hobson, G.M., et al.; Mutations in noncoding regions of the protelolipid protein gene in pelizaeous - Merzbacher disease, Neurology (2000) Vol. 55, pp. 1089-1096.	
	CL	Aradhya, Swaroop, et al.; Multiple pathogenic and benign genomic rearrangements occur at a 35 kb duplication involving the NEMO and LAGE2 genes, Hum. Mol. Genet. (2001), Vol. 10, No. 22, pp. 2557-2567.	
	CM	Galgoczy, Petra, et al.; Human-mouse comparative sequence analysis of the NEMO gene reveals an alternative promoter within the neighboring G6PD gene, Int'l Journal of Genes and Genomoes, (2001) Vol. 271, pp. 93-98.	

25384276.1 Examiner		Date Considered	3/16/04

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